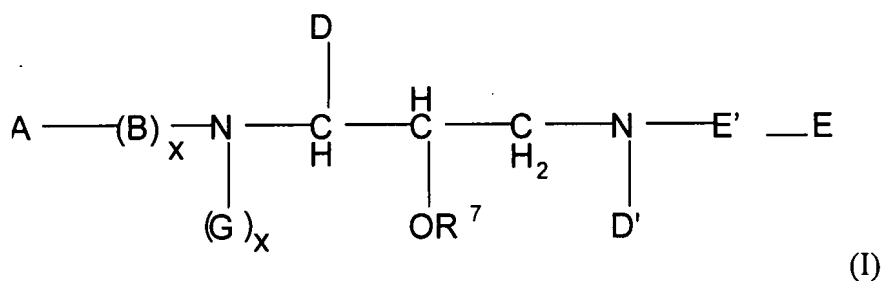


Amendments to the Claims

This listing of claims will replace all prior versions, and listings, of claims in the application:

1. (Currently amended) A compound of formula I:



or a pharmaceutically acceptable salt thereof, wherein:

E' is $\text{--SO}_2\text{--}$;

A is selected from $\text{--R}^1\text{--C}_1\text{--C}_6$ alkyl, which is optionally substituted with one or more groups independently selected from hydroxy, $\text{C}_1\text{--C}_4$ alkoxy, Ht, --O--Ht , $\text{--NR}^2\text{--CO--N(R}^2\text{)}_2$, $\text{--SO}_2\text{--R}^2$ or $\text{--CO--N(R}^2\text{)}_2$; or $\text{--R}^1\text{--C}_2\text{--C}_6$ alkenyl, which is optionally substituted with one or more groups independently selected from hydroxy, $\text{C}_1\text{--C}_4$ alkoxy, Ht, --O--Ht , $\text{--NR}^2\text{--CO--N(R}^2\text{)}_2$ or $\text{--CO--N(R}^2\text{)}_2$; or R^7 ;

R^1 is --O--C(O)-- ;

each Ht is independently selected from $\text{C}_3\text{--C}_7$ cycloalkyl; $\text{C}_5\text{--C}_7$ cycloalkenyl; $\text{C}_6\text{--C}_{14}$ aryl; or a 5-7 membered saturated or unsaturated heterocycle, containing one or more heteroatoms selected from N, O, or S; wherein said aryl or said heterocycle is optionally fused to Q; and wherein any member of said Ht is optionally substituted with one or more substituents independently selected from oxo, --OR^2 , SR^2 , --R^2 , $\text{--N(R}^2\text{)(R}^2\text{)}$, $\text{--R}^2\text{--OH}$, --CN , $\text{--CO}_2\text{R}^2$, $\text{--C(O)--N(R}^2\text{)}_2$, $\text{--S(O)}_2\text{--N(R}^2\text{)}_2$, $\text{--N(R}^2\text{)--C(O)--R}^2$, $\text{--N(R}^2\text{)--C(O)O--R}^2$, --C(O)--R^2 , $\text{--S(O)}_n\text{--}$

R^2 , $-OCF_3$, $-S(O)_n-Q$, methylenedioxy, $-N(R^2)-S(O)_2(R^2)$, halo, $-CF_3$, $-NO_2$, Q , $-OQ$, $-OR^7$, $-SR^7$, $-R^7$, $-N(R^2)(R^7)$ or $-N(R^7)_2$;

each Q is independently selected from a 3-7 membered saturated, partially saturated or unsaturated carbocyclic ring system; or a 5-7 membered saturated, partially saturated or unsaturated heterocyclic ring containing one or more heteroatoms selected from O, N, or S; wherein Q is optionally substituted with one or more groups selected from oxo, $-OR^2$, $-R^2$, $-SO_2R^2$, $-SO_2-N(R^2)_2$, $-N(R^2)_2$, $-N(R^2)-C(O)-R^2$, $-R^2-OH$, $-CN$, $-CO_2R^2$, $-C(O)-N(R^2)_2$, halo, $-CF_3$;

each R^2 is independently selected from H, or C_1-C_4 alkyl; and wherein said alkyl, when not a substituent of Q , is optionally substituted with Q or $-OR^3$; wherein when said R^2 is an $-OR^3$ substituted moiety, said R^3 in $-OR^3$ may not be $-OR^2$ substituted;

B is absent;

each x is independently 0 or 1;

each R^3 is independently selected from H, Ht, C_1-C_6 alkyl, C_2-C_6 alkenyl, C_2-C_6 alkynyl, C_3-C_6 cycloalkyl or C_5-C_6 cycloalkenyl; wherein any member of said R^3 , except H, is optionally substituted with one or more substituents selected from $-OR^2$, $-C(O)-NH-R^2$, $-S(O)_n-N(R^2)(R^2)$, $-N(R^2)_2$, $-N(R^2)-C(O)-O(R^2)$, $-N(R^2)-C(O)-N(R^2)$, $-N(R^2)-C(O)-(R^2)$, Ht, $-CN$, $-SR^2$, $-CO_2R^2$, or $NR^2-C(O)-R^2$;

each n is independently 1 or 2;

G is H;

D is **benzyl** ~~C_4-C_6 alkyl optionally substituted with Q~~ ;

D' is selected from C_1-C_{15} alkyl, C_2-C_{15} alkenyl or C_2-C_{15} alkynyl, each of which contains one or more substituents selected from oxo, $-CF_3$, $-OCF_3$, $-NO_2$, azido, $-SH$, $-N(R^3)-N(R^3)_2$, $-O-N(R^3)_2$, $-(R^3)N-O-(R^3)$, $-CN$, $-CO_2R^3$, $-C(O)-N(R^3)_2$, $-S(O)_n-N(R^3)_2$, $-N(R^3)-C(O)-R^3$, $-N(R^3)-C(O)-N(R^3)_2$, $-N(R^3)-C(O)-S(R^3)$, $-C(O)-R^3$, $-N(R^3)-S(O)_n(R^3)$, $-N(R^3)-S(O)_n-N(R^3)_2$, $-S-NR^3-C(O)R^3$, $-C(S)N(R^3)_2$, $-C(S)R^3$, $-NR^3-C(O)OR^3$, $-O-C(O)OR^3$, $-O-C(O)N(R^3)_2$,

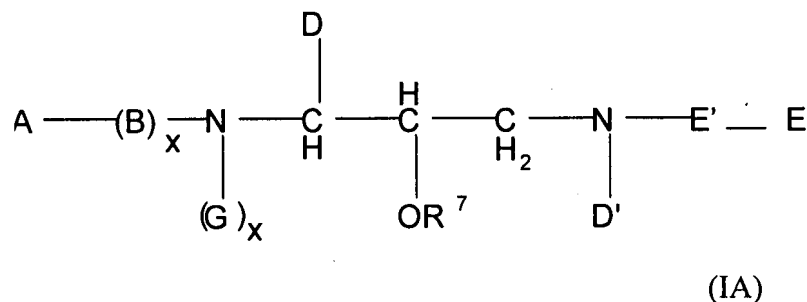
-NR³-C(S)R³, =N-OH, =N-OR³, =N-N(R³)₂, =NR³, =NNR³C(O)N(R³)₂, =NNR³C(O)OR³, =NNR³S(O)_n-N(R³)₂, -NR³-C(S)OR³, -NR³-C(S)N(R³)₂, -NR³-C[=N(R³)]-N(R³)₂, -N(R³)-C[=N-NO₂]-N(R³)₂, -N(R³)-C[=N-NO₂]-OR³, -N(R³)-C[=N-CN]-OR³, -N(R³)-C[=N-CN]-(R³)₂, -OC(O)R³, -OC(S)R³, -OC(O)N(R³)₂, -C(O)N(R³)-N(R³)₂, -O-C(O)N(R³)-N(R³)₂, O-C(O)N(OR³)(R³), N(R³)-N(R³)C(O)R³, N(R³)-OC(O)R³, N(R³)-OC(O)R³, N(R³)-OC(O)R³, -OC(S)N(R³)₂, -OC(S)N(R³)(R³), or PO₃-R³;

E is selected from Ht; O-Ht; Ht-Ht; Ht fused with Ht; -O-R³; -N(R²)(R³); C₁-C₆ alkyl optionally substituted with one or more groups selected from R⁴ or Ht; C₂-C₆ alkenyl optionally substituted with one or more groups selected from R⁴ or Ht; C₃-C₆ saturated carbocycle optionally substituted with one or more groups selected from R⁴ or Ht; or C₅-C₆ unsaturated carbocycle optionally substituted with one or more groups selected from R⁴ or Ht;

each R⁴ is independently selected from -OR², -OR³, -SR², -SOR², -SO₂R², -CO₂R², -C(O)-NHR², -C(O)-N(R²)₂, -C(O)-NR²(OR²), -S(O)₂-NHR², halo, -NR²-C(O)-R², -N(R²)₂ or -CN; and

each R⁷ is hydrogen.

2. (Previously presented) The compound according to claim 1, having the formula IA:



wherein:

D' is selected from C₁₋₁₅ alkyl, C₂₋₁₅ alkenyl or C₂.C₁₅ alkynyl; each of which is substituted with one to two –CN groups and is optionally substituted with C₃.C₈ cycloalkyl.

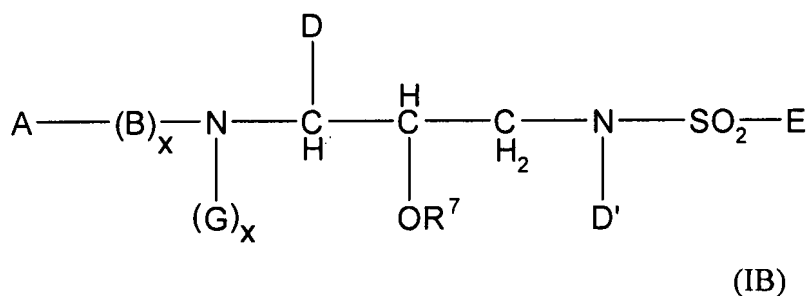
3. (Previously presented) The compound according to claim 2 wherein:

D' is selected from C₁₋₁₅ alkyl or C₂₋₁₅ alkenyl; each of which is substituted with one to two –CN groups and is optionally substituted with C₃.C₈ cycloalkyl.

4. (Previously presented) The compound according to claim 2 wherein:

D' is C₂.C₁₅ alkynyl which is substituted with one to two –CN groups and is optionally substituted with C₃.C₈ cycloalkyl.

5. (Previously presented) The compound according to claim 1 having the formula IB:



wherein:

D' is selected from C₁.C₁₅ alkyl, C₂.C₁₅ alkenyl or C₂.C₁₅ alkynyl, each of which contains one or more substituents selected from oxo, –CF₃, –OCF₃, –NO₂, azido, –SH, –N(R³)–N(R³)₂, –O–N(R³)₂, –(R³)N–O–(R³), –CO₂R³, –C(O)–N(R³)₂, –S(O)_n–N(R³)₂, –N(R³)–C(O)–R³, –N(R³)–C(O)–N(R³)₂, –N(R³)–C(O)–S(R³), –C(O)–R³, –N(R³)–S(O)_n(R³), –N(R³)–S(O)_n–N(R³)₂, –S–NR³–C(O)R³, –C(S)N(R³)₂, –C(S)R³, –NR³–C(O)OR³, –O–C(O)OR³, –O–C(O)N(R³)₂, –NR³–

$C(S)R^3$, $=N-OH$, $=N-OR^3$, $=N-N(R^3)_2$, $=NR^3$, $=NNR^3C(O)N(R^3)_2$, $=NNR^3C(O)OR^3$,
 $=NNR^3S(O)_n-N(R^3)_2$, $-NR^3-C(S)OR^3$, $-NR^3-C(S)N(R^3)_2$, $-NR^3-C[=N(R^3)]-N(R^3)_2$, $-N(R^3)-$
 $C[=N-NO_2]-N(R^3)_2$, $-N(R^3)-C[=N-NO_2]-OR^3$, $-N(R^3)-C[=N-CN]-OR^3$, $-N(R^3)-C[=N-CN]-$
 $(R^3)_2$, $-OC(O)R^3$, $-OC(S)R^3$, $-OC(O)N(R^3)_2$, $-C(O)N(R^3)-N(R^3)_2$, $-O-C(O)N(R^3)-N(R^3)_2$, $O-$
 $C(O)N(OR^3)(R^3)$, $N(R^3)-N(R^3)C(O)R^3$, $N(R^3)-OC(O)R^3$, $N(R^3)-OC(O)R^3$, $N(R^3)-OC(O)R^3$, $-$
 $OC(S)N(R^3)_2$, $-OC(S)N(R^3)(R^3)$, or PO_3-R^3 .

6. (Previously presented) The compound according to claim 5 wherein:

D' is selected from $C_1.C_{15}$ alkyl or $C_2.C_{15}$ alkenyl, each of which contains one or more
 substituents selected from oxo, $-CF_3$, $-OCF_3$, $-NO_2$, azido, $-N(R^3)-N(R^3)_2$, $-O-N(R^3)_2$, $-(R^3)N-$
 $O-(R^3)$, $-N(R^3)-C(O)-N(R^3)_2$, $-N(R^3)-C(O)-S(R^3)$, $-C(O)-R^3$, $-N(R^3)-S(O)_n(R^3)$, $-N(R^3)-S(O)_n-$
 $N(R^3)_2$, $-S-NR^3-C(O)R^3$, $-C(S)N(R^3)_2$, $-C(S)R^3$, $-NR^3-C(O)OR^3$, $-O-C(O)OR^3$, $-O-C(O)N(R^3)_2$,
 $-NR^3-C(S)R^3$, $=N-OH$, $=N-OR^3$, $=N-N(R^3)_2$, $=NR^3$, $=NNR^3C(O)N(R^3)_2$, $=NNR^3C(O)OR^3$,
 $=NNR^3S(O)_n-N(R^3)_2$, $-NR^3-C(S)OR^3$, $-NR^3-C(S)N(R^3)_2$, $-NR^3-C[=N(R^3)]-N(R^3)_2$, $-N(R^3)-$
 $C[=N-NO_2]-N(R^3)_2$, $-N(R^3)-C[=N-NO_2]-OR^3$, $-N(R^3)-C[=N-CN]-OR^3$, $-N(R^3)-C[=N-CN]-$
 $(R^3)_2$, $-OC(O)R^3$, $-OC(S)R^3$, $-OC(O)N(R^3)_2$, $-C(O)N(R^3)-N(R^3)_2$, $-O-C(O)N(R^3)-N(R^3)_2$, $O-$
 $C(O)N(OR^3)(R^3)$, $N(R^3)-N(R^3)C(O)R^3$, $N(R^3)-OC(O)R^3$, $N(R^3)-OC(O)R^3$, $N(R^3)-OC(O)R^3$,
 $-OC(S)N(R^3)_2$, $-OC(S)N(R^3)(R^3)$, or PO_3-R^3 ; $C_2.C_{15}$ alkynyl which contains one or more
 substituents selected from oxo, $-CF_3$, $-OCF_3$, $-NO_2$, azido, $-SH$, $-N(R^3)-N(R^3)_2$, $-O-N(R^3)_2$,
 $-(R^3)N-O-(R^3)$, $-CO_2R^3$, $-C(O)-N(R^3)_2$, $-S(O)_n-N(R^3)_2$, $-N(R^3)-C(O)-R^3$, $-N(R^3)-C(O)-N(R^3)_2$,
 $-N(R^3)-C(O)-S(R^3)$, $-C(O)-R^3$, $-N(R^3)-S(O)_n(R^3)$, $-N(R^3)-S(O)_n-N(R^3)_2$, $-S-NR^3-C(O)R^3$,
 $-C(S)N(R^3)_2$, $-C(S)R^3$, $-NR^3-C(O)OR^3$, $-O-C(O)OR^3$, $-O-C(O)N(R^3)_2$, $-NR^3-C(S)R^3$, $=N-OH$,
 $=N-OR^3$, $=N-N(R^3)_2$, $=NR^3$, $=NNR^3C(O)N(R^3)_2$, $=NNR^3C(O)OR^3$, $=NNR^3S(O)_n-N(R^3)_2$, $-NR^3-$
 $C(S)OR^3$, $-NR^3-C(S)N(R^3)_2$, $-NR^3-C[=N(R^3)]-N(R^3)_2$, $-N(R^3)-C[=N-NO_2]-N(R^3)_2$, $-N(R^3)-$
 $C[=N-NO_2]-OR^3$, $-N(R^3)-C[=N-CN]-OR^3$, $-N(R^3)-C[=N-CN]-N(R^3)_2$, $-OC(O)R^3$, $-OC(S)R^3$, $-$
 $OC(O)N(R^3)_2$, $-C(O)N(R^3)-N(R^3)_2$, $-O-C(O)N(R^3)-N(R^3)_2$, $O-C(O)N(OR^3)(R^3)$, $N(R^3)-$
 $N(R^3)C(O)R^3$, $N(R^3)-OC(O)R^3$, $N(R^3)-OC(O)R^3$, $N(R^3)-OC(O)R^3$, $-OC(S)N(R^3)_2$,
 $-OC(S)N(R^3)(R^3)$, or PO_3-R^3 .

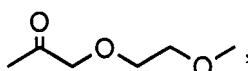
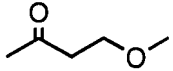
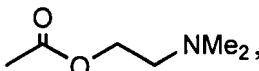
7. (Previously presented) The compound according to claim 5 wherein:

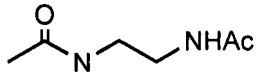
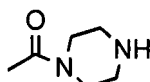
D' is selected from C₁-C₁₅ alkyl or C₂-C₁₅ alkenyl, each of which contains one or more substituents selected from -SH, -CO₂R³, -C(O)-N(R³)₂, -S(O)_n-N(R³)₂ or -N(R³)-C(O)-R³.

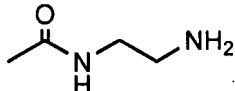
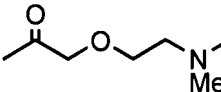
8. (Canceled)

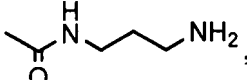
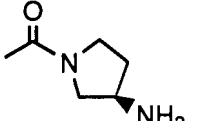
9. (Withdrawn – Currently amended) The compound according to any one

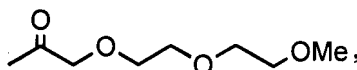
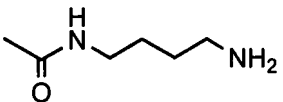
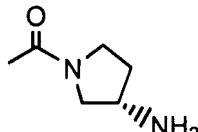
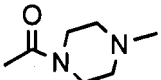
of claims 1 to 7, wherein at least one R⁷ is selected from:


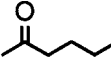
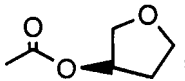
 ,  , -(L)-lysine, -PO₃Na₂,  ,

 , -(L)-tyrosine,  , -PO₃Mg, -PO₃(NH₄)₂, -CH₂-OPO₃Na₂,

 , -(L)-serine, -SO₃Na₂,  , -SO₃Mg, -SO₃(NH₄)₂ ,

CH₂-OSO₃Na₂, -CH₂-OSO₃(NH₄)₂,  ,  ,

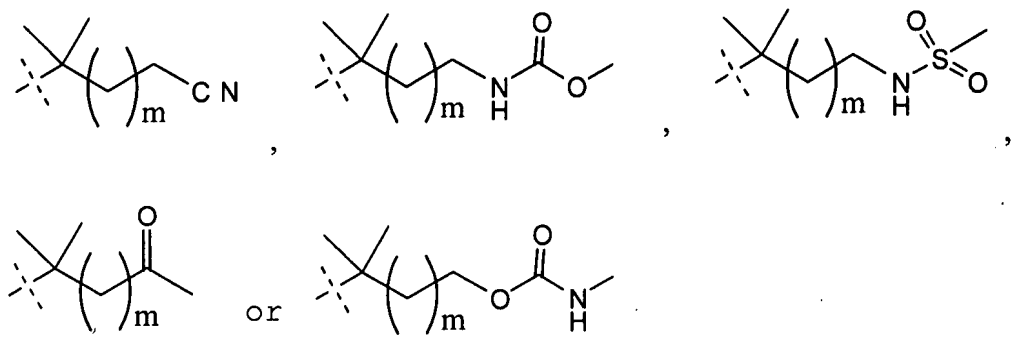
 ,  ,  ,  , acetyl,

 ,  , -(L)-valine, -(L)-glutamic acid, -(L)-aspartic acid, -(L)-γ-t-butyl-aspartic acid,  ,

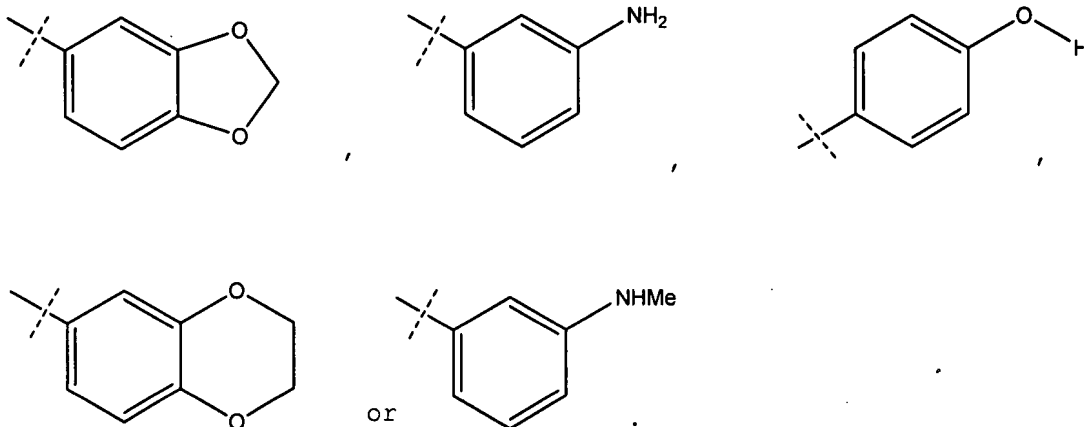
CC(=O)OC1CCOC1, CC(=O)OCCCCO[C@H]1[C@@H](OC(=O)C)[C@H](OC(=O)C)[C@H](OC(=O)C)[C@H](O)[C@H]1O, [O-]P(=O)([O-])OCCN,
[O-]P(=O)([O-])OCCN, CCOP(=O)([O-])[O-], CCOS(=O)([O-])[O-], PO3K2, PO3Ca,

10-11. (Canceled)

R'' is selected from

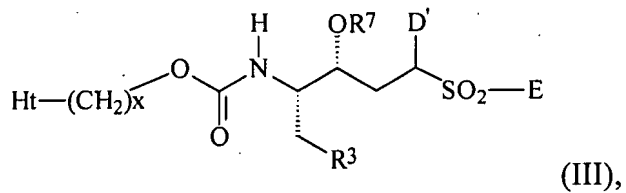


13. (Original) The compound according to claim 10, wherein E is selected



14. (Withdrawn) The compound according to claim 10, wherein R^7 is $-\text{PO}_3^{2-}$

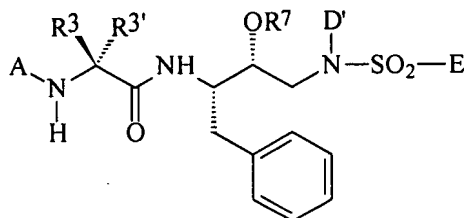
15. (Currently amended) The compound according to claim 1, having the formula III:



wherein $x = 1$; **and**

R^3 is phenyl.

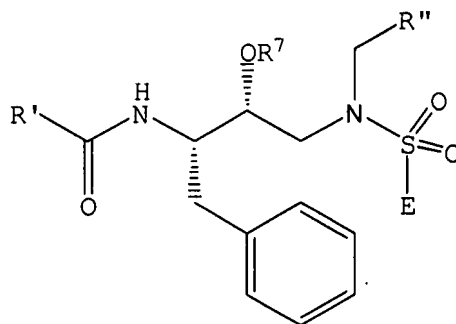
16. (Withdrawn) The compound according to claim 1, having the formula IV:



(IV);

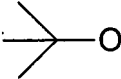
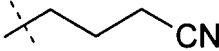
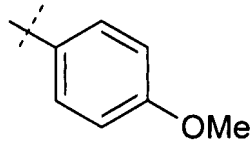
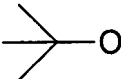
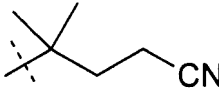
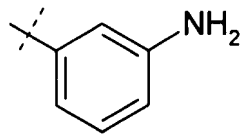
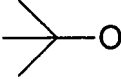
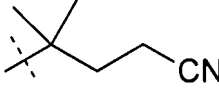
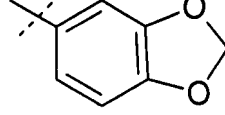
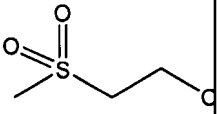
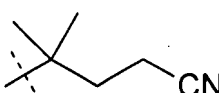
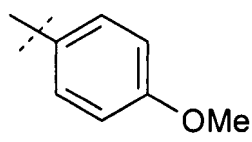
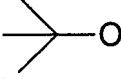
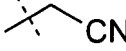
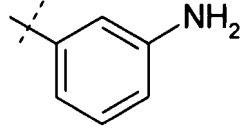
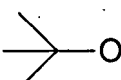
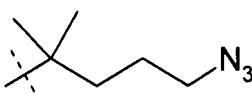
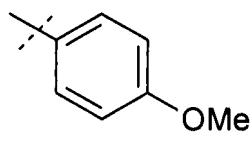
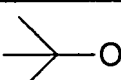
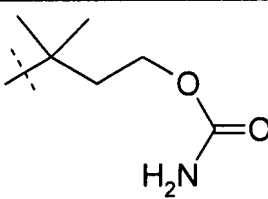
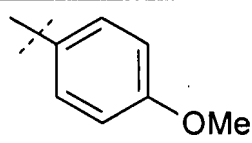
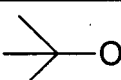
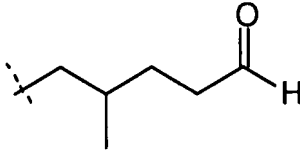
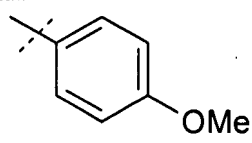
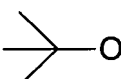
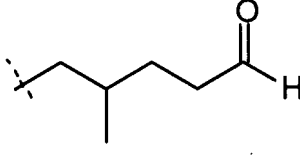
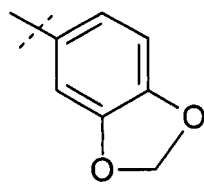
wherein $R^{3'}$ is selected from H, Ht, C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_3 - C_6 cycloalkyl or C_5 - C_6 cycloalkenyl; wherein any member of said R^3 , except H, is optionally substituted with one or more substituents selected from $-OR^2$, $-C(O)-NH-R^2$, $-S(O)_n-N(R^2)(R^2)$, $-N(R^2)_2$, $-N(R^2)-C(O)-O(R^2)$, $-N(R^2)-C(O)-N(R^2)$, $-N(R^2)-C(O)-(R^2)$, $-N(R^2-OR^2)_2$, $-C(O)-Ht$, Ht, $-CN$, $-SR^2$, $-CO_2R^2$, or $NR^2-C(O)-R^2$.

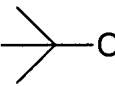
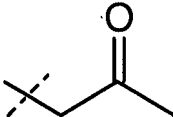
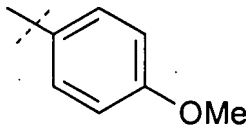
17. (Currently amended) The compound according to claim 1, wherein said compound is selected from any one of compound numbers: 1, 2, 3, 4, 5, 6, 22, 127, 203, 234, 277, 278, **and** 279, ~~363~~, and 364:

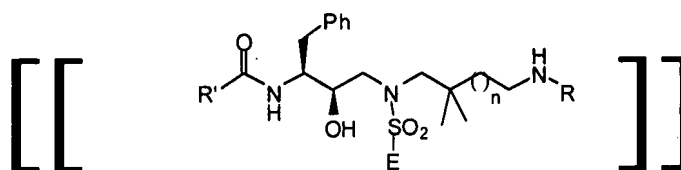


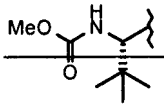
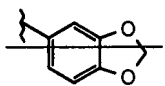
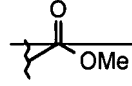
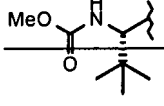
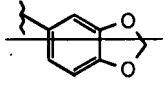
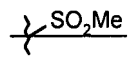
wherein R^7 is H; and

Compound	R'	R''	E
1			
2			
3			

Compound	R'	R''	E
4			
5			
6			
22			
127			
203			
234			
277			
278			

Compound	R'	R''	E
279			



Compound	R'	E	n	R
363			3	
364			3	

18-22. (Canceled)

23. (Currently amended) A composition comprising a compound according to any one of claims 1-7, [[10,]] 12, 13, 15, and 17 or a pharmaceutically acceptable salt thereof in a therapeutically effective amount, and a pharmaceutically acceptable carrier.

24. (Canceled)

25. (Original) The composition according to claim 23, wherein said composition is formulated as a pharmaceutically acceptable, orally available tablet or capsule.

26. (Previously presented) A method of treating an HIV virus infection in a human comprising the step of administering to said human a composition according to claim 23.

27-29. (Canceled)

30. (Previously presented) A method of treating an HIV virus infection in a human comprising the step of administering to said human a composition according to claim 25.

31. (Canceled)